ABSTRACT OF THE DISCLOSURE

The invention relates to a compound of formula (I)

or a salt thereof: wherein: R¹ is C₁₋₄alkyl, C₁₋₃fluoroalkyl or -(CH₂)₂OH; R² is a hydrogen atom (H), methyl or C₁fluoroalkyl; R^{3a} is a hydrogen atom (H) or C₁₋₃alkyl; R³ is optionally substituted branched C₃₋₆alkyl, optionally substituted C₃₋₈cycloalkyl, optionally substituted mono-unsaturated-C₅₋₇cycloalkenyl, optionally substituted phenyl, or an optionally substituted heterocyclic group of sub-formula (aa), (bb) or (cc): in which n¹ and n² independently are 1 or 2; and Y is O, S, SO₂, or NR⁴; and wherein Het is of sub-formula (i), (ii), (iii), (iv) or (v): The compounds are phosphodiesterase (PDE) inhibitors, in particular PDE4 inhibitors. Also provided is the use of a compound of formula (I), or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment and/or prophylaxis of an inflammatory and/or allergic disease in a mammal such as a human, for example chronic obstructive pulmonary disease (COPD), asthma, or allergic rhinitis.